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OY 256 TGCTGAGGCGCGCATGTTCTCTCACCCTTCCCCCGCGCGCTGTTAGAGAAAGTCT 315
    |||||
DB 241 TGCTGAGCGCGCATGTTCTCTCACCCTTCCCCCGCGCGCTGTTAGAGAAAGTCT 300
OY 316 TCCAGTACATTGACCTCCATCAGAGTGAATTTGTGCAGACGCTGAAGGAGTGGTGCCA 375
    |||||
DB 301 TCCAGTACATTGACCTCCATCAGAGTGAATTTGTGCAGACGCTGAAGGAGTGGTGCCA 360
OY 376 TCGAGAGGCACTCTGTCCAGCCTGTGCTCTCAGACAAGAGCTCTTCAGAAATGATGG 435
    |||||
DB 361 TCGAGAGGCACTCTGTCCAGCCTGTGCTCTCAGACAAGAGCTCTTCAGAAATGATGG 420
OY 436 CCGTGCTGCGGGAACACGCTGCAGCGCTGGGGGGCCGCTGTGGCTCGGTGACATGGGTC 495
    |||||
DB 421 CCGTGCTGCGGGAACACGCTGCAGCGCTGGGGGGCCGCTGTGGCTCGGTGACATGGGTC 480
OY 496 CTCAGCAGCTGCCCGATGGTCAAGTCTTCCAATACCTCCGCTCATCTGGCCGAACTGG 555
    |||||
DB 481 CTCAGCAGCTGCCCGATGGTCAAGTCTTCCAATACCTCCGCTCATCTGGCCGAACTGG 540
OY 556 GGAGCGATCCCAAGAAAGGACCGTGTGCTTCTACGGCCAATTGGAAGTGCAGCTGCTG 615
    |||||
DB 541 GGAGCGATCCCAAGAAAGGACCGTGTGCTTCTACGGCCAATTGGAAGTGCAGCTGCTG 600
OY 616 ACCGGGGGATGGGTGGCTCAACGACCCCTATGTGCTGACGGAAGTAGACGGGAACTTT 675
    |||||
DB 601 ACCGGGGGATGGGTGGCTCAACGACCCCTATGTGCTGACGGAAGTAGACGGGAACTTT 660
OY 676 ATGACGAGAGGACGACCGACAACAAAGGCCCTGTGCTGGCTGGATCAATGCTGTGAGCG 735
    |||||
DB 661 ATGACGAGAGGACGACCGACAACAAAGGCCCTGTGCTGGCTGGATCAATGCTGTGAGCG 720
OY 736 CCTTCAGAGCCCTGAGAGAGATCTTCTGTGAATATCAAAATTCATCATGAGGGATGG 795
    |||||
DB 721 CCTTCAGAGCCCTGAGAGAGATCTTCTGTGAATATCAAAATTCATCATGAGGGATGG 780
OY 796 AAGAGCTGGCTCTGTGCTCTGAGAGAACTGTGAGAAAGAAAGAGACCGATTCTTCT 855
    |||||
DB 781 AAGAGCTGGCTCTGTGCTCTGAGAGAACTGTGAGAAAGAAAGAGACCGATTCTTCT 840
OY 856 CTGTGTGAGACTACATTTGAATTTTCAATTAACCTGTGATCAGCCAAAGAACCCAGCAA 915
    |||||
DB 841 CTGTGTGAGACTACATTTGAATTTTCAATTAACCTGTGATCAGCCAAAGAACCCAGCAA 900
OY 916 TCACCTAATGAAACCCGGGGGAAACAGCTACTTCATGTGAGGTGAATGACAGACCAAG 975
    |||||
DB 901 TCACCTAATGAAACCCGGGGGAAACAGCTACTTCATGTGAGGTGAATGACAGACCAAG 960
OY 976 ATTTTCACTAGGAACCTTGGTGGCATCTTCATGAACCAATGCTGATCTGTGCTC 1035
    |||||
DB 961 ATTTTCACTAGGAACCTTGGTGGCATCTTCATGAACCAATGCTGATCTGTGCTC 1020
OY 1036 TTCTCGGTAGCCTGTGAGACTCGTGTGCTATATCCTGTGCCCTGGAATCTATGATGAAG 1095
    |||||
DB 1021 TTCTCGGTAGCCTGTGAGACTCGTGTGCTATATCCTGTGCCCTGGAATCTATGATGAAG 1080
OY 1096 TGGTCTCTTACAGAGAGAAATAATACATACAAAGCCATCCATCTAGACCTAGAAG 1155
    |||||
DB 1081 TGGTCTCTTACAGAGAGAAATAATACATACAAAGCCATCCATCTAGACCTAGAAG 1140
OY 1156 AATACCGGAATAGCAGCGGGGTGAGAAATTTCTGTTCGATACTAAGAGAGAGATTCTAA 1215
    |||||
DB 1141 AATACCGGAATAGCAGCGGGGTGAGAAATTTCTGTTCGATACTAAGAGAGAGATTCTAA 1200
OY 1216 TGCACCTCTGAGGTACCATCTCTTTCTAATTCATGGATCGAGGGCGGCTTGATGAGC 1275
    |||||
DB 1201 TGCACCTCTGAGGTACCATCTCTTTCTAATTCATGGATCGAGGGCGGCTTGATGAGC 1260
OY 1276 CTGGAATAAAACAGTCATACCTGCGGAGTTATAGAAATTTTCAATCCGCTACTGCC 1335
    |||||
DB 1261 CTGGAATAAAACAGTCATACCTGCGGAGTTATAGAAATTTTCAATCCGCTACTGCC 1320
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OY 1336 CTCACATGAATGTGTCTGCGGTGGAATAACAGAGTGACACGACATCTTGAAGATGTCTCT 1395
    |||||
DB 1321 CTCACATGAATGTGTCTGCGGTGGAATAACAGAGTGACACGACATCTTGAAGATGTCTCT 1380
OY 1396 CCAAAAGAAATAGTTCACACAGATGGTGTTCCTAGTACTCTAGGACTACACCCGTGGA 1455
    |||||
DB 1381 CCAAAAGAAATAGTTCACACAGATGGTGTTCCTAGTACTCTAGGACTACACCCGTGGA 1440
OY 1456 TTGCAATATTGATGACACCCAGTATCTCGCAGCAAAAAGAGCGATCAGAACAGTGTTCG 1515
    |||||
DB 1441 TTGCAATATTGATGACACCCAGTATCTCGCAGCAAAAAGAGCGATCAGAACAGTGTTCG 1500
OY 1516 GAACAGAACAGATATGATCCGGGATGGATCCACCATTCCTCAATTGCCCCAAATGTTCCAGG 1575
    |||||
DB 1501 GAACAGAACAGATATGATCCGGGATGGATCCACCATTCCTCAATTGCCCCAAATGTTCCAGG 1560
OY 1576 AGATCGTCCACAAGAGCGGTGTGCTAATTCGCTGGAGCTGTGATGATGAGAAACATT 1635
    |||||
DB 1561 AGATCGTCCACAAGAGCGGTGTGCTAATTCGCTGGAGCTGTGATGATGAGAAACATT 1620
OY 1636 CGCAGAAATGAGAAATCAACAGGTGGAACATAAGAGGAAACCAATTAATTGCTGCCCT 1695
    |||||
DB 1621 CGCAGAAATGAGAAATCAACAGGTGGAACATAAGAGGAAACCAATTAATTGCTGCCCT 1680
OY 1696 TTTTCTTAGAGATGGCCAGCTCCATTAATCAAGAACCTTCTAGTCTGATCTGATCCCA 1755
    |||||
DB 1681 TTTTCTTAGAGATGGCCAGCTCCATTAATCAAGAACCTTCTAGTCTGATCTGATCCCA 1740
OY 1756 CTGACAGATTACCTCCCAACATCCCTAGACAGGAGTGAATGTAAATATCCAGAGAAAT 1815
    |||||
DB 1741 CTGACAGATTACCTCCCAACATCCCTAGACAGGAGTGAATGTAAATATCCAGAGAAAT 1800
OY 1816 TTGGGTCTAGTATAGTACATTTTCCCTTCCATTTAAATGCTTGGGATATCTGATCAG 1875
    |||||
DB 1801 TTGGGTCTAGTATAGTACATTTTCCCTTCCATTTAAATGCTTGGGATATCTGATCAG 1860
OY 1876 TAAATAAATATTTCAAAAGGCACAGATGTTGAATGTTAAAGTCCCCCACTGCACACC 1935
    |||||
DB 1861 TAAATAAATATTTCAAAAGGCACAGATGTTGAATGTTAAAGTCCCCCACTGCACACC 1920
OY 1936 TTCTCAAGTACATAGCTGCTTGACGAACAATTGTTCCCAAGTCCGTGTGCAATAGCCCC 1995
    |||||
DB 1921 TTCTCAAGTACATAGCTGCTTGACGAACAATTGTTCCCAAGTCCGTGTGCAATAGCCCC 1980
OY 1996 AGGATTGGATTCCTTCCAACTTTTGAATATCTCCAACTTGCAATTTGATGGCATAA 2055
    |||||
DB 1981 AGGATTGGATTCCTTCCAACTTTTGAATATCTCCAACTTGCAATTTGATGGCATAA 2040
OY 2056 TCACTCCGGTTGCTTCTTACGCTCAAGTCTCGTGACACATAATCATTCATCCAAT 2115
    |||||
DB 2041 TCACTCCGGTTGCTTCTTACGCTCAAGTCTCGTGACACATAATCATTCATCCAAT 2100
OY 2116 GATCGCTTTGCTTTACCACTCTTCTTATCTTATTAATAAAATGTTG 2167
    |||||
DB 2101 GATCGCTTTGCTTTACCACTCTTCTTATCTTATTAATAAAATGTTG 2152
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RESULT 2
AA10638/c
ID AA10638 standard, DNA; 127 BP.

XX AA10638;

DT 30-MAR-1999 (first entry)

DE Human biallelic polymorphic DNA fragment WI-15225.

KW Polymorphism; biallelic; human; forensic; paternity testing; disease;

KW detection; phenotypic typing; characteristic; infection; hereditary;

KW autoimmune disease; cancer; inflammation; drug; therapy; medication;

OS Homo sapiens.

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XX PN MO9820165-A2.
XX PD 14-MAY-1998.
XX PF 05-NOV-1997; 97WO-US020313.
XX PR 06-NOV-1996; 96US-0030455P.
XX PA (WHED ) WHITEHEAD INST BIOMEDICAL RES.
XX PI Lander ES, Wang D, Hudson T;
XX DR WPI; 1998-286974/25.
XX PT New isolated nucleic acid segments from the human genome - used for
XX PT determining polymorphic forms for use in e.g. forensics, paternity
XX PT testing or phenotypic typing for disease.
XX PS Claim 1; Page 67; 310pp; English.
XX CC AAX10269-X12937 are human DNA fragments which contain biallelic
XX CC polymorphic markers which have been isolated using the primers
XX CC represented in AAX09121-X10268. The base occupying the polymorphic site
XX CC is indicated by the appropriate IUPAC-IUB ambiguity code. These fragments
XX CC can be used in methods for determining polymorphic forms in an individual
XX CC for use in e.g. forensics, paternity testing or for phenotypic typing for
XX CC diseases such as agammaglobulinemia, diabetes insipidus, Lesch-Nyhan
XX CC syndrome, muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease,
XX CC familial hypercholesterolemia, polycystic kidney disease, hereditary
XX CC spherocytosis, von Willebrand's disease, tuberous sclerosis, hereditary
XX CC haemorrhagic telangiectasia, familial colonic polyposis, Ehlers-Danlos
XX CC syndrome, osteogenesis imperfecta, acute intermittent porphyria,
XX CC autoimmune diseases, inflammation, cancer, diseases of the nervous
XX CC system, infection by pathogenic microorganisms, and characteristics such
XX CC as longevity, appearance (e.g. baldness, obesity), strength, speed,
XX CC endurance, fertility, and susceptibility or receptivity to particular
XX CC drugs or therapeutic treatments. The isolated polymorphic nucleic acid
XX CC segments can also be used to produce medicaments for the treatment or
XX CC prophylaxis of such diseases
XX SQ Sequence 127 BP; 47 A; 16 C; 36 G; 27 T; 0 U; 1 Other;

Query Match 5.6%; Score 126.6; DB 1; Length 127;
Best Local Similarity 99.2%; Pred. No. 9.2;
Matches 126; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2017 TTTTAGCATATCTCCAACTTGCAATTGATGGCATATACCTCCGCTTGTCTTAG 2076
DB 127 TTTTAGCATATCTCCAACTTGCAATTGATGGCATATACCTCCGCTTGTCTTAG 68

QY 2077 GTCTCAAGTGCTCGTGACACATATATTCATTCCAATGATGCGCTTGTCTTAGCACT 2136
DB 67 GTCTCAAGTGCTCGTGACACATATATTCATTCCAATGATGCGCTTGTCTTAGCACT 8

QY 2137 CTTTCTCT 2143
DB 7 CTTTCTCT 1

RESULT 3
AAV81394
ID AAV81394 standard; DNA; 1733 BP.
XX AC AAV81394;
XX DE 16-MAR-1999 (first entry)
XX DE Human tumour antigen zsig15 coding sequence.
XX KM Secretion; differentiation marker; tumour; epithelial cell; colon; blood;
XX KM breast; prostate; growth; development; antagonist; receptor; bone marrow;
XX KM cancer; metastasis; ss.

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XX OS Homo sapiens.
XX FH Key Location/Qualifiers
XX FT CDS 34..1347
XX FT /*tag= a
XX FT /product= "zsig15"
XX PN WO9850552-A1.
XX PD 12-NOV-1998.
XX PF 06-MAY-1998; 98WO-US009584.
XX PR 06-MAY-1997; 97US-0045703P.
XX PA (ZYMO ) ZYMOGENETICS INC.
XX PI Shepard PO, Grossmann A;
XX DR WPI; 1999-034723/03.
XX DR P-PSDB; AAW67722.
XX PT New nucleic acid encoding secreted polypeptide zsig15 - used as a marker
XX PT for tumour cells, useful for diagnosis and treatment of cancers,
XX PT inflammation and hyperplasia.
XX PS Claim 4; Page 81-84; 100pp; English.
XX CC This sequence encodes a secreted polypeptide, designated zsig15, which is
XX CC a marker for differentiation in normal and tumour cells (particularly
XX CC epithelial cells and derived tumours of colon, breast and prostate). The
XX CC zsig15 protein is useful for the promotion of growth and development of
XX CC epithelial cells; to identify specific (ant)agonists, also where
XX CC conjugated to a toxin, to deliver these to cells expressing the cognate
XX CC receptor (e.g. to kill cells of blood, colon, breast and bone marrow
XX CC cancers), and to identify/isolate receptors involved in cancer
XX CC metastases. The sequence was isolated from a colon cancer library after
XX CC screening an EST (expressed sequence tag) database for sequences
XX SQ Sequence 1733 BP; 448 A; 504 C; 442 G; 339 T; 0 U; 0 Other;

Query Match 2.8%; Score 62.4; DB 1; Length 1733;
Best Local Similarity 96.9%; Pred. No. 1.3;
Matches 63; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2178 CTGNCCTCCCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 2237
DB 1599 CTTTCTCCCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 1658

QY 2238 AAAAA 2242
DB 1659 AAAAA 1663

RESULT 4
AAZ65270/C
ID AAZ65270 standard; DNA; 2152 BP.
XX AC AAZ65270;
XX DE 23-MAR-2000 (first entry)
XX DE Human secreted protein gene 21.
XX KM Human; secreted protein; cancer; tumour; developmental abnormality;
XX KM foetal deficiency; blood disorder; immune system disorder; inflammation;
XX KM autoimmune disease; allergy; Alzheimer's disease; cognitive disorder;
XX KM schizophrenia; arthritis; asthma; psoriasis; sepsis; skin disorder;
XX KM atherosclerosis; diabetes; cardiovascular disorder; kidney disorder;
XX KM digestive disorder; endocrine disorder; infection; AIDS; leukaemia;
XX KM therapy; chromosome 18q22-23; ds.

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